



# House of Representatives

General Assembly

**File No. 101**

February Session, 2012

Substitute House Bill No. 5329

*House of Representatives, March 26, 2012*

The Committee on General Law reported through REP. TABORSAK of the 109th Dist., Chairperson of the Committee on the part of the House, that the substitute bill ought to pass.

## **AN ACT CONCERNING THE USE OF TELEPHARMACY BY HOSPITALS.**

Be it enacted by the Senate and House of Representatives in General Assembly convened:

1 Section 1. Section 50 of public act 11-242 is repealed and the  
2 following is substituted in lieu thereof (*Effective July 1, 2012*):

3 (a) As used in this section:

4 (1) "Electronic technology" or "telepharmacy" means the process: (A)  
5 By which each step involved in the [preparation of IV admixtures]  
6 dispensing of a sterile product is verified through use of a bar code  
7 tracking system and documented by means of digital photographs  
8 which are electronically recorded and preserved; and (B) which is  
9 monitored and verified through video and audio communication  
10 between a licensed supervising [clinical] pharmacist and a pharmacy  
11 technician;

12 (2) ["IV admixture" means an IV fluid to which one or more

13 additional drug products have been added] "Sterile product" means  
14 any drug, as that term is defined in section 20-571 of the general  
15 statutes, that is compounded, manipulated or otherwise prepared  
16 under sterile conditions during the dispensing process, is not intended  
17 for self-administration by a patient and is intended to be used in a  
18 hospital, or its satellite, remote or affiliated office-based locations;

19 (3) "Pharmacist" means an individual who is licensed to practice  
20 pharmacy under the provisions of section 20-590, 20-591, 20-592 or  
21 20-593 of the general statutes, and who is thereby recognized as a  
22 health care provider by the state of Connecticut; and

23 (4) "Pharmacy technician" means an individual who is registered  
24 with the department and qualified in accordance with section 20-598a  
25 of the general statutes.

26 (b) [The Commissioner of Consumer Protection, in consultation  
27 with the Commissioner of Public Health, may establish a pilot  
28 program to permit a] A hospital, licensed in accordance with the  
29 provisions of chapter 368v of the general statutes, which operates a  
30 hospital pharmacy, [to] may use electronic technology or telepharmacy  
31 at the hospital and at the hospital's satellite or remote locations for  
32 purposes of allowing a [clinical] pharmacist to supervise pharmacy  
33 technicians in the [preparation of IV admixtures] dispensing of sterile  
34 products. [Under the pilot program, notwithstanding]  
35 Notwithstanding the provisions of chapter 400j of the general statutes  
36 or regulations adopted pursuant to said chapter, a [clinical] pharmacist  
37 shall be permitted to supervise a pharmacy technician through use of  
38 electronic technology, [A supervising clinical] and under such  
39 supervision the pharmacist shall monitor and verify the activities of a  
40 pharmacy technician through audio and video communication. The  
41 pharmacist-to-technician ratio pursuant to section 20-576-33 of the  
42 regulations of Connecticut state agencies shall apply. In the event of a  
43 malfunction of the electronic technology, no [IV admixtures] sterile  
44 product prepared by a pharmacy technician during the time period of  
45 the malfunction may be distributed to patients, unless [an

46 appropriately licensed individual] a licensed pharmacist is able to: (1)  
47 Personally review and verify the accuracy of all processes utilized in  
48 the [preparation of the IV admixture] dispensing of the sterile product;  
49 or (2) upon the restoration of the electronic technology, utilize the  
50 mechanisms of the electronic technology which recorded the actions of  
51 the pharmacy technician to confirm that all proper steps were followed  
52 in the [preparation of the IV admixture] dispensing of the sterile  
53 product. [Under the pilot program, all] All orders for [medication]  
54 sterile products to be dispensed using telepharmacy shall be verified  
55 by a pharmacist prior to being delegated to a pharmacy technician for  
56 [preparation of an IV admixture] such dispensing. A hospital  
57 [participating in the pilot program] shall ensure that appropriately  
58 licensed personnel administer medications [at the hospital's satellite or  
59 remote locations] dispensed using telepharmacy. All of the processes  
60 involved in [the operation of the pilot program] a hospital's use of  
61 telepharmacy shall be under the purview of the hospital's director of  
62 pharmacy.

63 (c) A hospital [selected to participate in the pilot program] using  
64 telepharmacy shall undertake periodic quality assurance evaluations,  
65 not less than once per calendar quarter, which shall [minimally]  
66 include, upon discovery, prompt review of any error in medication  
67 administration which occurs [under the pilot program] where  
68 telepharmacy is used to dispense such medication. A hospital shall  
69 make such quality assurance evaluations available for review and  
70 inspection by the Departments of Consumer Protection and Public  
71 Health.

72 [(d) A pilot program established pursuant to this section may  
73 commence operation on or after July 1, 2011, and shall terminate not  
74 later than December 31, 2012, provided the Commissioner of  
75 Consumer Protection may terminate the pilot program prior to  
76 December 31, 2012, for good cause shown.]

This act shall take effect as follows and shall amend the following sections:
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Section 1	<i>July 1, 2012</i>	PA 11-242, Sec. 50
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**GL**      *Joint Favorable Subst.*

The following Fiscal Impact Statement and Bill Analysis are prepared for the benefit of the members of the General Assembly, solely for purposes of information, summarization and explanation and do not represent the intent of the General Assembly or either chamber thereof for any purpose. In general, fiscal impacts are based upon a variety of informational sources, including the analyst's professional knowledge. Whenever applicable, agency data is consulted as part of the analysis, however final products do not necessarily reflect an assessment from any specific department.

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***OFA Fiscal Note***

***State Impact:*** None

***Municipal Impact:*** None

***Explanation***

There is no fiscal impact to the Department of Consumer Protection (DCP) making the telepharmacy program permanent as the DCP had successfully piloted the program and already has the resources in place.

***The Out Years***

***State Impact:*** None

***Municipal Impact:*** None

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**OLR Bill Analysis****sHB 5329*****AN ACT CONCERNING THE USE OF TELEPHARMACY BY HOSPITALS.*****SUMMARY:**

This bill makes permanent the telepharmacy pilot program and expands it to (1) all licensed hospital pharmacies and (2) dispense sterile products, not just IV admixture preparations as under the pilot program. It allows pharmacists at hospital pharmacies to use electronic technology at the hospital, its satellite, or remote locations to allow a clinical pharmacist to supervise pharmacy technicians in dispensing sterile products.

Under the bill, the pilot program ends on July 1, 2012.

EFFECTIVE DATE: July 1, 2012

**TELEPHARMACY**

The bill expands the application of electronic technology or telepharmacy to dispensing sterile products. The pilot program limits the use of this technology to preparing IV admixtures, which also involves sterile products.

Under the pilot program and the bill, “electronic technology” or “telepharmacy” means the process (1) by which each step involved in the dispensing of sterile products is verified by a bar code tracking system and documented by digital photographs that are electronically recorded and preserved and (2) which is monitored and verified through video and audio communication between a licensed supervising clinical pharmacist and a pharmacy technician.

**STERILE PRODUCTS**

Sterile products are any drug that is compounded, manipulated, or otherwise prepared under sterile conditions during the dispensing process. It is not intended for self-administration by a patient and is intended to be used in a hospital, its satellite, remote, or affiliated office-based location. Under the pilot program, technicians could only dispense IV admixtures, which is an IV fluid to which one or more additional drug products have been added.

## **PROGRAM REQUIREMENTS**

Under the bill, a pharmacist is authorized to supervise a pharmacy technician dispensing sterile products through electronic technology and monitor and verify the technician's activities through audio and video communication. The number of technicians the pharmacist can supervise must conform to the existing regulatory pharmacy-to-technician ratio. For inpatient and satellite pharmacies, that ratio is 3:1, which can be increased to 5:1 on the pharmacy director's petition and Pharmacy Commission approval.

The bill applies the current procedures for electronic technology malfunctions involving IV admixtures to those involving any sterile product. If the electronic technology malfunctions, no sterile product prepared by the pharmacy technician during the malfunction period can be distributed to patients unless a licensed pharmacist can (1) personally review and verify all of the processes used in preparing the sterile product or (2) after the technology is restored, use the electronic technology mechanisms that recorded the pharmacy technician's actions to confirm that all proper steps were followed in preparing the sterile product. All orders for medication must be verified by a pharmacist before being delegated to a pharmacy technician for sterile product preparation.

As with the pilot program, the bill requires a hospital to ensure that appropriately licensed health care personnel administer medications at the hospital's satellite or remote locations. The bill specifies that all processes involved in operating the program are under the purview of the hospital's pharmacy director.

**EVALUATIONS**

The bill extends the current requirement for periodic quality assurance to telepharmacies used to dispense sterile products. It specifically requires hospitals to make periodic quality assurance evaluations, at least once per calendar quarter, which includes, upon discovery, prompt review of any error in medication administration. The hospital must make these evaluations available to the departments of Consumer Protection and Public Health for their review.

**COMMITTEE ACTION**

General Law Committee

Joint Favorable Substitute

Yea 17 Nay 0 (03/13/2012)